What is claimed is :

1. An embryonic stem cell line derived from a nucleus-transferred oocyte prepared by transferring a nucleus of a human somatic cell into an enucleated human oocyte.

2. The embryonic stem cell line of claim 1, which is a cell line deposited under the accession number of KCLRF-BP-00092.

3. A method for preparing an embryonic stem cell line, comprising the steps of:

(1) culturing a human somatic cell to prepare a nuclear donor cell;

(2) enucleating a human oocyte to prepare a recipient oocyte;

(3) preparing a nucleus-transferred oocyte by transferring a nucleus of the nuclear donor cell into the recipient oocyte and fusing the nucleus of the nuclear donor cell and the recipient oocyte;

(4) subjecting the nucleus-transferred oocyte to reprogramming, activation and *in vitro* culturing to form a blastocyst; and

(5) isolating an inner cell mass from the blastocyst and culturing the inner cell mass in an undifferentiated state to establish the embryonic stem cell line.

4. The method of claim 3, wherein the embryonic stem cell line is a cell line deposited under the accession number of KCLRF-BP-00092.

5. The method of claim 3, wherein the reprogramming in step (4) is conducted for a time period of up to 20 hours.

6. The method of claim 3, wherein the reprogramming in step (4) is conducted for a time period of up to 6 hours.

7. The method of claim 3, wherein the reprogramming in step (4) is conducted for a time period of up to 3 hours.

8. The method of claim 3, wherein the reprogramming in step (4) is conducted for a time period of about 2 hours.

9. The method of claim 3, wherein the activation in step (4) is performed by treating the nucleus-transferred oocyte with a calcium ionophore and subsequently with 6-dimethylaminopurine.

10. The method of claim 9, wherein the concentration of the calcium ionophore ranges from 5μM to 15μM.

11. The method of claim 9, wherein the concentration of the calcium ionophore is about 10μM.

12. The method of claim 9, wherein the concentration of 6-dimethylaminopurine ranges from 1.5mM to 2.5mM.

13. The method of claim 9, wherein the concentration of 6-dimethylaminopurine is about 2.0mM.

14. The method of claim 3, wherein the *in vitro* culturing in step (4) is performed by sequentially using at least two media, each having a different composition from the other.

15. The method of claim 14, wherein the *in vitro* culturing is performed by sequentially using two media having different compositions each other.

16. The method of claim 15, wherein the *in vitro* culturing is performed by sequentially using the G1.2 medium and the SNUnt-2 medium.

17. The method of claim 3, wherein step (4) is performed by reprogramming the nucleus-transferred oocyte for a time period of up to 20 hours, treating the nucleus-transferred oocyte with a calcium ionophore at a concentration ranging from 5μM to 15μM and subsequently with 6-dimethylaminopurine at a concentration ranging from 1.5mM to 2.5mM, and sequentially culturing the nucleus-transferred oocyte *in vitro* in the G1.2 medium and the SNUnt-2 medium.

18. The method of claim 3, wherein the inner cell mass is isolated from the blastocyst in step (5) by a process comprising the steps of:

(1) removing the zona pellucida or part thereof from the blastocyst; and

(2) isolating the inner cell mass by removing the trophoblast from the resulting blastocyst.

19. The method of claim 3, wherein the inner cell mass is cultured in step (5) on a feeder layer comprising a cell differentiated from the embryonic stem cell line of claim 1.

20. A neuro progenitor differentiated from an embryonic stem cell line derived from a nucleus-transferred oocyte prepared by transferring a nucleus of a human somatic cell into an enucleated human oocyte.

21. The neuro progenitor of claim 20, wherein the embryonic stem cell line is a cell line deposited under the accession number of KCLRF-BP-00092.

22. A method for preparing the neuro progenitor of claim 20, comprising the steps of:

(1) culturing the embryonic stem cell line to form an embryoid body;

(2) culturing the embryoid body in the presence of an agent suitable for differentiating a cell of the embryoid body into the neuro progenitor; and

(3) selecting a cell expressing a marker of the neuro progenitor and culturing the selected cell to obtain the neuro progenitor.

23. The method of claim 22, wherein the embryonic stem cell line is a cell line deposited under the accession number of KCLRF-BP-00092.

24. The method of claim 22, wherein the agent employed in step (2) is selected from the group consisting of retinoic acid; ascorbic acid; nicotinamide; N-2 supplement; B-27 supplement; and a mixture of insulin, transferrin, sodium selenite and fibronectin.

25. A medium for use in carrying out the *in vitro* culturing in step (4) of claim 3, comprising:

95 to 110mM NaCl; 7.0 to 7.5mM KCl; 20 to 30mM $NaHCO_3$,; 1.0 to 1.5mM NaH_2PO_4; 3 to 8mM sodium lactate; 1.5 to 2.0mM $CaCl_2 \cdot 2H_2O$; 0.3 to 0.8mM $MgCl_2 \cdot 6H_2O$; 0.2 to 0.4mM sodium pyruvate; 1.2 to 1.7mM fructose; 6 to 10㎎/㎖ human serum albumin; 0.7 to 0.8㎍/㎖ kanamycin; 1.5 to 3% essential amino acids; 0.5 to 1.5% nonessential amino acids; 0.7 to 1.2mM L-glutamine; and 0.3 to 0.7% a mixture of insulin, transferrin and sodium selenite.

26. The medium of claim 25, which comprises:

99.1 to 106mM NaCl; 7.2mM KCl; 25mM $NaHCO_3$; 1.2mM NaH_2PO_4; 5mM sodium lactate; 1.7mM $CaCl_2 \cdot 2H_2O$; 0.5mM $MgCl_2 \cdot 6H_2O$; 0.3mM sodium pyruvate; 1.5mM fructose; 8㎎/㎖ human serum albumin; 0.75㎍/㎖ kanamycin; 2% essential amino acids; 1% nonessential amino acids; 1mM L-glutamine; and 0.5% a mixture of insulin, transferrin and sodium selenite.